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Date Mailed:	April	_1_	_, 2003	

Sheet 1 of 2 Allarlie to # 25

FORM LE JANORMATION DISCLOSURE STATEMENT	Docket Number: 9192.5USWO	Application Number: 09 117,071	
IN AN APPLICATION	Applicant: Kingsman et al.		
(Use several sheets if necessary)	Filing Date: 09/25/1998	Group Art Unit: 1636	

			l	U.S. PATENT DOCUME	NTS			
EXAMINER INITIAL	DOCUMENT NO.  US 6,503,501 B1		. DATE NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE		
52			01/07/03	Anderson et al.	6.261	43 2	5/13/99	
			FOI	REIGN PATENT DOCU	MENTS			
	DOCUMENT NO.		DATE COUNTR	COUNTRY	CLASS	SUBCLASS	TRANSLATION	
							YES	NO
		OTHER	DOCUMENT	S (Including Author, Title.	Date Pertinent I	Pages Ftc.)		
	1					ages, Lie.)		
5{	Brigham et al., 2000, <i>Hum. Gene Ther.</i> , 11:1023-32  Transfection of nasal mucosa with a normal alpha 1-antitrypsin gene in alpha1-antitrypsin-deficient subjects: comparison with protein therapy							
St.	_	Caplen et al., 1995, Nat. Med., 1:39-46  Liposome-mediated CFTR gene transfer to the nasal epithelium of patients with cystic fibrosis						
Se ,		Gleich et al., 1998, Arch Otolaryngol Head Neck Surg., 124:1097-104  Alloantigen gene therapy for squamous cell carcinoma of the head and neck: results of a phase-1 trial						
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Successful F		ful Peripheral T-Lymphocyte-Directed Gene Transfer for a Patient With Severe Combined Immune ney Caused by Adenosine Deaminase Deficiency						

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EXAMINER	1. 1. 1. 1.	DATE CONSIDERED	(1/3/02

EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form for next communication to the Applicant.

09 117,071

INFORMATION DISCLOSURE STATEMENT

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FORM 1/ 🏈

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Group Art Unit: 1636

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TEM & TRA	(	Porteous et al., 1997, Gene Ther., 4:210-218
Si	,	Evidence for safety and efficacy of DOTAP cationic liposome mediated CFTR gene transfer to the nasal epithelium of patients with cystic fibrosis
×	7.	Raper et al., 1996, <i>Annals of Surgery</i> , 223:116-126 Safety and Feasibility of Liver-Directed <i>Ex Vivo</i> Gene Therapy for homozygous Familial Hypercholesterolemia
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